RVD2008-04

Re-evaluation Decision

Triallate

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Re-evaluation Decision

After a re-evaluation of the herbicide triallate, Health Canada's Pest Management Regulatory Agency (PMRA), under the authority of the *Pest Control Products Act* and Regulations, is granting continued registration of certain triallate products for sale and use in Canada.

An evaluation of available scientific information found that, under the proposed conditions of use, products containing triallate do not present unacceptable risks to human health or the environment when used according to label directions. As a condition of the continued registration of triallate uses, new risk-reduction measures must be included on the label of products containing triallate. Additional data are required to maintain specific uses.

The major risk-mitigation measures are as follows:

- For emulsifiable concentrate formulations, label statements are required to limit the amount handled per day to 189 kg active ingredient, and to indicate that closed mix/load systems and application with a closed cab are required.
- Additional personal protective equipment (PPE) is required for granular formulations.
- A restricted-entry interval (REI) of 12 hours is required to enter or allow worker entry into treated areas.
- Additional precautionary measures to prevent runoff, leaching and volatization are required.
- Terrestrial and aquatic buffer zones for emulsifiable concentrate formulations are required.

The regulatory approach regarding the re-evaluation of triallate was first proposed in the Proposed Re-evaluation Decision document PRVD2007-08, *Re-evaluation of Triallate*, a consultation document.[1] This Re-evaluation Decision document[2] describes this stage of the PMRA's regulatory process concerning the re-evaluation of triallate and summarizes the Agency's decision and the reasons for it. No comments were received during the consultation process. Therefore, this decision has not changed from the proposal stated in PRVD2007-08. To comply with this decision, registrants of triallate products will be informed of the specific requirements affecting their product registration(s) and of the regulatory options available to them.

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1 "Consultation statement" as required by subsection 28(2) of the *Pest Control Products Act*.

2 "Decision statement" as required by subsection 28(5) of the *Pest Control Products Act*.

For more details on the information presented in this Re-evaluation Decision, please refer to the related Proposed Re-evaluation Decision, PRVD2007-08, *Re-evaluation of Triallate*.

Other Information

For triallate, the summary of assessments found in PRVD2007-08 serves as an evaluation report. A list of references considered by the Agency in support of the registration decision is found in this Re-evaluation Decision document (Appendix I). The relevant test data on which the decision is based are available for public inspection, upon application, in the PMRA's Reading Room (located in Ottawa). For more information, please contact the PMRA's Pest Management Information Service by phone (1-800-267-6315) or by e-mail (pmra_infoserv@hc-sc.gc.ca).

Any person may file a notice of objection[3] regarding this decision on triallate within 60 days from the date of publication of this Re-evaluation Decision document. For more information regarding the basis for objecting (which must be based on scientific grounds), please refer to the PMRA's website (Requesting a Reconsideration of Decision, www.pmra-arla.gc.ca/english/pubreg/reconsideration-e.html), or contact the PMRA's Pest Management Information Service by phone (1-800-267-6315) or by e-mail (pmra_infoserv@hc-sc.gc.ca).

[3] As per subsection 35(1) of the *Pest Control Products Act*

Appendix I List of References

A. Studies/Information Provided by the Applicant/Registrant (Unpublished)

Impact on Human and Animal Health

PMRA Document Number	Reference
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